# PDGFRA-associated chronic eosinophilic leukemia

*PDGFRA*-associated chronic eosinophilic leukemia is a form of blood cell cancer characterized by an elevated number of cells called eosinophils in the blood. These cells help fight infections by certain parasites and are involved in the inflammation associated with allergic reactions. However, these circumstances do not account for the increased number of eosinophils in *PDGFRA*-associated chronic eosinophilic leukemia.

Another characteristic feature of *PDGFRA*-associated chronic eosinophilic leukemia is organ damage caused by the excess eosinophils. Eosinophils release substances to aid in the immune response, but the release of excessive amounts of these substances causes damage to one or more organs, most commonly the heart, skin, lungs, or nervous system. Eosinophil-associated organ damage can lead to a heart condition known as eosinophilic endomyocardial disease, skin rashes, coughing, difficulty breathing, swelling (edema) in the lower limbs, confusion, changes in behavior, or impaired movement or sensations. People with *PDGFRA*-associated chronic eosinophilic leukemia can also have an enlarged spleen (splenomegaly) and elevated levels of certain chemicals called vitamin B12 and tryptase in the blood.

Some people with *PDGFRA*-associated chronic eosinophilic leukemia have an increased number of other types of white blood cells, such as neutrophils or mast cells. Occasionally, people with *PDGFRA*-associated chronic eosinophilic leukemia develop other blood cell cancers, such as acute myeloid leukemia or B-cell or T-cell acute lymphoblastic leukemia or lymphoblastic lymphoma.

*PDGFRA*-associated chronic eosinophilic leukemia is often grouped with a related condition called hypereosinophilic syndrome. These two conditions have very similar signs and symptoms; however, the cause of hypereosinophilic syndrome is unknown.

## Frequency

*PDGFRA*-associated chronic eosinophilic leukemia is a rare condition; however, the exact prevalence is unknown.

# **Genetic Changes**

PDGFRA-associated chronic eosinophilic leukemia is caused by mutations in the PDGFRA gene. This condition usually occurs as a result of genetic rearrangements that fuse part of the PDGFRA gene with part of another gene. Rarely, changes in single DNA building blocks (point mutations) in the PDGFRA gene are found in people with this condition. Genetic rearrangements and point mutations affecting the PDGFRA gene are somatic mutations, which are mutations acquired during a person's lifetime that are present only in certain cells. The somatic mutation occurs initially in a single cell,

which continues to grow and divide, producing a group of cells with the same mutation (a clonal population).

The most common genetic abnormality in *PDGFRA*-associated chronic eosinophilic leukemia results from a deletion of genetic material from chromosome 4, which brings together part of the *PDGFRA* gene and part of the *FIP1L1* gene, creating the *FIP1L1-PDGFRA* fusion gene.

The *FIP1L1* gene provides instructions for a protein that plays a role in forming the genetic blueprints for making proteins (messenger RNA or mRNA).

The *PDGFRA* gene provides instructions for making a receptor protein that is found in the cell membrane of certain cell types. Receptor proteins have specific sites into which certain other proteins, called ligands, fit like keys into locks. When the ligand attaches (binds), the PDGFRA receptor protein is turned on (activated), which leads to activation of a series of proteins in multiple signaling pathways. These signaling pathways control many important cellular processes, such as cell growth and division (proliferation) and cell survival.

The FIP1L1-PDGFRA fusion gene (as well as other PDGFRA fusion genes) provides instructions for making a fusion protein that has the function of the normal PDGFRA protein. However, the fusion protein does not require ligand binding to be activated. Similarly, point mutations in the PDGFRA gene can result in a PDGFRA protein that is activated without ligand binding. As a result, the signaling pathways are constantly turned on (constitutively activated), which increases the proliferation and survival of cells. When the FIP1L1-PDGFRA fusion gene mutation or point mutations in the PDGFRA gene occur in blood cell precursors, the growth of eosinophils (and occasionally other blood cells, such as neutrophils and mast cells) is poorly controlled, leading to PDGFRA-associated chronic eosinophilic leukemia. It is unclear why eosinophils are preferentially affected by this genetic change.

#### Inheritance Pattern

PDGFRA-associated chronic eosinophilic leukemia is not inherited and occurs in people with no history of the condition in their families. Mutations that lead to a PDGFRA fusion gene and PDGFRA point mutations are somatic mutations, which means they occur during a person's lifetime and are found only in certain cells. Somatic mutations are not inherited. Males are more likely to develop PDGFRA-associated chronic eosinophilic leukemia than females because, for unknown reasons, PDGFRA fusion genes are found more often in males.

#### Other Names for This Condition

PDGFRA-associated myeloproliferative neoplasm

# **Diagnosis & Management**

# **Genetic Testing**

 Genetic Testing Registry: Idiopathic hypereosinophilic syndrome https://www.ncbi.nlm.nih.gov/gtr/conditions/C0206141/

## Other Diagnosis and Management Resources

- Cancer.Net: Leukemia Eosinophilic: Treatment http://www.cancer.net/cancer-types/leukemia-eosinophilic/treatment-options?se ctionTitle=Treatment
- MedlinePlus Encyclopedia: Eosinophil Count Absolute https://medlineplus.gov/ency/article/003649.htm
- Seattle Cancer Care Alliance: Hypereosinophilia https://www.seattlecca.org/diseases/blood-disorders/eosinophilic-disorders

### General Information from MedlinePlus

- Diagnostic Tests
   https://medlineplus.gov/diagnostictests.html
- Drug Therapy https://medlineplus.gov/drugtherapy.html
- Genetic Counseling https://medlineplus.gov/geneticcounseling.html
- Palliative Care https://medlineplus.gov/palliativecare.html
- Surgery and Rehabilitation https://medlineplus.gov/surgeryandrehabilitation.html

#### Additional Information & Resources

## MedlinePlus

- Encyclopedia: Eosinophil Count Absolute https://medlineplus.gov/ency/article/003649.htm
- Health Topic: Eosinophilic Disorders https://medlineplus.gov/eosinophilicdisorders.html

## Genetic and Rare Diseases Information Center

 Hypereosinophilic syndrome https://rarediseases.info.nih.gov/diseases/2804/hypereosinophilic-syndrome

### Additional NIH Resources

 National Cancer Institute: Chronic Eosinophilic Leukemia https://www.cancer.gov/types/myeloproliferative/patient/chronic-treatment-pdq#section/\_258

### **Educational Resources**

- Cancer.Net: Leukemia Eosinophilic http://www.cancer.net/cancer-types/leukemia-eosinophilic?sectionTitle=Overview
- Cancer.Net: Leukemia Eosinophilic: Treatment http://www.cancer.net/cancer-types/leukemia-eosinophilic/treatment-options?se ctionTitle=Treatment
- Cincinnati Children's: What is an Eosinophil? https://www.cincinnatichildrens.org/service/c/eosinophilic-disorders/conditions/eosinophil
- Disease InfoSearch: Idiopathic hypereosinophilic syndrome http://www.diseaseinfosearch.org/Idiopathic+hypereosinophilic+syndrome/3753
- MalaCards: pdgfra-associated chronic eosinophilic leukemia http://www.malacards.org/card/pdgfra\_associated\_chronic\_eosinophilic\_leukemia
- Merck Manual for Health Care Professionals: Eosinophilia http://www.merckmanuals.com/professional/hematology-and-oncology/eosinophilic-disorders/eosinophilia
- Orphanet: Hypereosinophilic syndrome http://www.orpha.net/consor/cgi-bin/OC\_Exp.php?Lng=EN&Expert=168956

# Patient Support and Advocacy Resources

- American Partnership for Eosinophilic Disorders http://apfed.org/
- Hanns A. Pielenz Clinical Research Center for Myeloproliferative Neoplasia, MD Anderson Cancer Center https://www.mdanderson.org/research/departments-labs-institutes/programs-centers/clinical-research-center-for-myeloproliferative-neoplasia.html

## ClinicalTrials.gov

ClinicalTrials.gov
 https://clinicaltrials.gov/ct2/results?cond=%22PDGFRA-associated+chronic
 +eosinophilic+leukemia%22+OR+%22Hypereosinophilic+Syndrome%22

#### Scientific Articles on PubMed

PubMed

https://www.ncbi.nlm.nih.gov/pubmed?term=%28chronic+eosinophilic+leukemia %5BTIAB%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1440+days%22%5Bdp%5D

#### **OMIM**

 HYPEREOSINOPHILIC SYNDROME, IDIOPATHIC http://omim.org/entry/607685

## **Sources for This Summary**

- Buitenhuis M, Verhagen LP, Cools J, Coffer PJ. Molecular mechanisms underlying FIP1L1-PDGFRA-mediated myeloproliferation. Cancer Res. 2007 Apr 15;67(8):3759-66.
   Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/17440089
- Cools J, DeAngelo DJ, Gotlib J, Stover EH, Legare RD, Cortes J, Kutok J, Clark J, Galinsky I, Griffin JD, Cross NC, Tefferi A, Malone J, Alam R, Schrier SL, Schmid J, Rose M, Vandenberghe P, Verhoef G, Boogaerts M, Wlodarska I, Kantarjian H, Marynen P, Coutre SE, Stone R, Gilliland DG. A tyrosine kinase created by fusion of the PDGFRA and FIP1L1 genes as a therapeutic target of imatinib in idiopathic hypereosinophilic syndrome. N Engl J Med. 2003 Mar 27;348(13):1201-14. Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/12660384
- Elling C, Erben P, Walz C, Frickenhaus M, Schemionek M, Stehling M, Serve H, Cross NC, Hochhaus A, Hofmann WK, Berdel WE, Müller-Tidow C, Reiter A, Koschmieder S. Novel imatinibsensitive PDGFRA-activating point mutations in hypereosinophilic syndrome induce growth factor independence and leukemia-like disease. Blood. 2011 Mar 10;117(10):2935-43. doi: 10.1182/blood-2010-05-286757. Epub 2011 Jan 11.
   Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/21224473
- Fukushima K, Matsumura I, Ezoe S, Tokunaga M, Yasumi M, Satoh Y, Shibayama H, Tanaka H, Iwama A, Kanakura Y. FIP1L1-PDGFRalpha imposes eosinophil lineage commitment on hematopoietic stem/progenitor cells. J Biol Chem. 2009 Mar 20;284(12):7719-32. doi: 10.1074/jbc.M807489200. Epub 2009 Jan 14.
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/19147501
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2658066/
- Metzgeroth G, Walz C, Score J, Siebert R, Schnittger S, Haferlach C, Popp H, Haferlach T, Erben P, Mix J, Müller MC, Beneke H, Müller L, Del Valle F, Aulitzky WE, Wittkowsky G, Schmitz N, Schulte C, Müller-Hermelink K, Hodges E, Whittaker SJ, Diecker F, Döhner H, Schuld P, Hehlmann R, Hochhaus A, Cross NC, Reiter A. Recurrent finding of the FIP1L1-PDGFRA fusion gene in eosinophilia-associated acute myeloid leukemia and lymphoblastic T-cell lymphoma. Leukemia. 2007 Jun;21(6):1183-8. Epub 2007 Mar 22.

Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/17377585

- Roufosse FE, Goldman M, Cogan E. Hypereosinophilic syndromes. Orphanet J Rare Dis. 2007 Sep 11;2:37. Review.
  - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/17848188
    Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2045078/
- Vandenberghe P, Wlodarska I, Michaux L, Zachée P, Boogaerts M, Vanstraelen D, Herregods MC, Van Hoof A, Selleslag D, Roufosse F, Maerevoet M, Verhoef G, Cools J, Gilliland DG, Hagemeijer A, Marynen P. Clinical and molecular features of FIP1L1-PDFGRA (+) chronic eosinophilic leukemias. Leukemia. 2004 Apr;18(4):734-42.
   Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/14973504

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